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CLAIMS

What is claimed is:

- 1. A pharmaceutical composition comprising a polypeptide comprising an HMGB B box or a functional variant thereof, in an amount sufficient to treat a disease or condition by increasing an immune response in an individual administered said pharmaceutical composition.
- 2. The pharmaceutical composition of Claim 1, wherein said HMGB B box is mammalian.
- 3. The pharmaceutical composition of Claim 2, wherein said HMGB B box is human.
 - 4. The pharmaceutical composition of Claim 3, wherein said polypeptide comprises an HMGB1 B box polypeptide.
 - 5. The pharmaceutical composition of Claim 4, wherein said polypeptide consists of an HMGB1 B box polypeptide.
- 15 6. The pharmaceutical composition of Claim 1, further comprising a vaccine.
 - 7. The pharmaceutical composition of Claim 6, further comprising an adjuvant.
 - 8. The pharmaceutical composition of Claim 7, wherein said adjuvant is selected from the group consisting of one or more immunostimulatory oligonucleotides, an imidazoquinoline, monophosphoryl lipid A, and detoxified lipopolysaccharide.

- The pharmaceutical composition of Claim 8, wherein said immunostimulatory oligonucleotides comprise unmethylated CpG sequences.
- 10. An antibody attached to a polypeptide comprising an HMGB B box or a functional variant thereof.
- 5 11. The antibody of Claim 10, wherein said HMGB B box is mammalian.
 - 12. The antibody of Claim 11, wherein said HMGB B box is human.
 - 13. The antibody of Claim 12, wherein said polypeptide comprises an HMGB1 B box polypeptide.
- The antibody of Claim 13, wherein said polypeptide consists of an HMGB1
 B box polypeptide.
 - 15. The antibody of Claim 10, wherein said antibody binds a tumor-associated polypeptide.
 - 16. The antibody of Claim 10, wherein said antibody is in a pharmaceutically acceptable carrier.
 - 17. A method of stimulating or increasing an immune response in an individual in need of immunostimulation, said method comprising administering to said individual a polypeptide comprising an HMGB B box or a functional variant thereof, in a amount sufficient to stimulate or increase said immune response.
- 20 18. The method of Claim 17, wherein said individual is being treated for cancer.
 - 19. The method of Claim 17, wherein said HMGB B box is mammalian.

- 20. The method of Claim 19, wherein said HMGB B box is human.
- 21. The method of Claim 20, wherein said polypeptide comprises an HMGB1 B box.
- 22. The method of Claim 21, wherein said polypeptide consists of an HMGB1 B box.
 - 23. The method of Claim 17, wherein said polypeptide is co-administered with a vaccine.
 - 24. The method of Claim 23, wherein said polypeptide is co-administered with a further adjuvant.
- 10 25. The method of Claim 24, wherein said adjuvant is selected from the group consisting of one or more immunostimulatory oligonucleotides, an imidazoquinoline, monophosphoryl lipid A, and detoxified lipopolysaccharide.
- 26. The method of Claim 25, wherein said immunostimulatory oligonucleotides comprise unmethylated CpG sequences.
 - 27. The method of Claim 17, wherein said administration is systemic.
 - 28. The method of Claim 17, wherein said administration is localized to a target site.
- 29. The method of Claim 17, wherein said polypeptide is attached to an antibody specific to a target site in the individual in need of immunostimulation.

- 30. The method of Claim 17, wherein said polypeptide is in a pharmaceutically acceptable carrier.
- 31. A method of treating cancer in an individual, said method comprising administering to said individual a therapeutically effective amount of a polypeptide comprising an HMGB B box or a functional variant thereof.
- 32. The method of Claim 31, wherein said HMGB B box is mammalian.
- 33. The method of Claim 32, wherein said HMGB B box is human.
- 34. The method of Claim 33, wherein said polypeptide comprises an HMGB1 B box polypeptide.
- 10 35. The method of Claim 34, wherein said polypeptide consists of an HMGB1 B box polypeptide.
 - 36. The method of Claim 31, wherein said polypeptide is co-administered with a vaccine.
- 37. The method of Claim 36, wherein said polypeptide is co-administered with a further adjuvant.
 - 38. The method of Claim 37, wherein said adjuvant is selected from the group consisting of one or more immunostimulatory oligonucleotides, an imidazoquinoline, monophosphoryl lipid A, and detoxified lipopolysaccharide.
- 20 39. The method of Claim 38, wherein said immunostimulatory oligonucleotides comprise unmethylated CpG sequences.

- 40. The method of Claim 31, wherein said administration is systemic.
- 41. The method of Claim 31, wherein said administration is localized to a target site.
- 42. The method of Claim 41, wherein said target site is a tumor.
- 5 43. The method of Claim 31, wherein said polypeptide is attached to an antibody.
 - 44. The method of Claim 43, wherein said antibody binds a tumor-associated polypeptide.
- The method of Claim 31, wherein said polypeptide is in a pharmaceutically acceptable carrier.